Under the Paperwork Reduction Act of 1995, no persons are required to re

INFORMATION DISCLOSURE
STATEMENT BY APPLICANT
( Not for submission under 37 CFR 1.99)

Application Number		10535294	
Filing Date		2002-05-17	
First Named Inventor	Elmo	M. A. Diederiks	
Art Unit		2821	
Examiner Name			
Attamen Dealest Numb		NII 024400	

				Attorney Docket Number NL021199			NL021199				
					U.S.	PATENTS				Remove	
Examiner Initial*	Cite No	Patent Number	Kind Code <sup>1</sup>	Issue E	Issue Date Name of Patentee or Applicant of cited Document		Pages,Columns,Lines where Relevant Passages or Relevant Figures Appear				
	1										
If you wish	h to a	i dd additional U.S. Pate	nt citatio	n inform	ation pl	lease click the	Ad	d button.		Add	
			U.S.P	ATENT	APPLI	CATION PUBL	LIC	ATIONS		Remove	
Examiner Initial*	Cite No	Publication Number	Kind Code <sup>1</sup>	Publica Date	tion	of cited Decument Releva		s,Columns,Lines when ant Passages or Rele es Appear			
	1	20030218537	A1	2003-1	1-27	HOCH ET A	NL.				
If you wish	h to a	dd additional U.S. Publi	shed Ap	plication	citatio	n information p	olea	se click the Ad	butto	n. Add	
				FOREIG	SN PAT	TENT DOCUM	ΕN	ITS		Remove	
Examiner Initial*	Cite No	Foreign Document Number <sup>3</sup>	Country Code <sup>2</sup>		Kind Code <sup>4</sup>	Publication Date	A	ame of Patented pplicant of cited ocument		Pages, Columns, Lines where Relevant Passages or Relevan Figures Appear	74
	1	JP2001060406			A	2001-03-06		MATSUSHITA L	то		
	2	JP2000036392			A	2000-02-02		TOSHIBA CORE	,		
If you wish	h to a	dd additional Foreign P				information pl			buttor	Add	_

	Application Number		10535294
INFORMATION DIGGS COURT	Filing Date		2002-05-17
INFORMATION DISCLOSURE STATEMENT BY APPLICANT	First Named Inventor	Elmo	M. A. Diederiks
(Not for submission under 37 CFR 1.99)	Art Unit		2821
(·	Examiner Name		
	Attorney Docket Numb	er	NL021199

Exan	niner (	Cite	Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposium, catalog, etc), date, pages(s), volume-issue number(s), publisher, city and/or country where published.	Ţ5	
		1			

If you wish to add additional non-patent literature document citation information please click the Add button Add

EXAMINER SIGNATURE

# Examiner Signature Date Considered

\*EXAMINER: Initial if reference considered, whether or not citation is in conformance with MPEP 609. Draw line through a citation if not in conformance and not considered. Include copy of this form with next communication to applicant.

1 See Knd Codes of USPTO Patent Documents at year LISPTO, GDU or MPEP 901.04. <sup>2</sup> Either office that issued the document, by the bo-claims code (WIPO Standard ST3.) <sup>3</sup> Sor Juapanes patent concuments, by managed the common state of the Empower managed code the search counterful. <sup>4</sup> Knill of document by the appropriate symbols as adicated on the document under WIPO Standard ST.16 if possible. <sup>3</sup> Applicant is to place a check mark here if English Imaguage translation is attached.

## INFORMATION DISCLOSURE STATEMENT BY APPLICANT (Not for submission under 37 CFR 1.99)

Application Number		10535294				
Filing Date		2002-05-17				
First Named Inventor	Elmo	M. A. Diederiks				
Art Unit		2821				
Examiner Name						
Attorney Docket Number		NL021199				

#### CERTIFICATION STATEMENT

Please see 37 CFR 1.97	and 1.98 to make the	appropriate selection(s):
------------------------	----------------------	---------------------------

That each item of information contained in the information disclosure statement was first cited in any communication from a foreign patent office in a counterpart foreign application not more than three months prior to the filing of the information siclosure statement. See 37 CFR 197(e)(1).

## OR

That no item of information contained in the information disclosure statement was cited in a communication from a foreign patent office is a counterpart foreign application, and, to the knowledge of the person signing the certification after making reasonable inquiry, no item of information contained in the information disclosure statement was known to any individual designated in 37 CFR 1/5(c) more than three months prior to the filing of the information disclosure statement. See 37 CFR 1/9/(c).

- See attached certification statement.
- Fee set forth in 37 CFR 1.17 (p) has been submitted herewith.
- 7 None

#### SIGNATURE

A signature of the applicant or representative is required in accordance with CFR 1.33, 10.18. Please see CFR 1.4(d) for the form of the signature.

_			
Signature	/Yan Glickberg/	Date (YYYY-MM-DD)	2006-10-04
Name/Print	Yan Glickberg	Registration Number	51742

This collection of information is required by 3T CFR 1.97 and 1.98. The information is required to obtain or retain a benefit by the public which is for life and by the USPTO to process) an application. Confidentiality is governed by \$5 U.S. C.12 and 3T CFR.

1.14. This collection is estimated to take I hour to complete, including gathering, preparing and submitting the completed application from the USPTO. Time will vary depending upon the individual case. Any comments on the amount of time you require to complete this form and/or suggestions for reducing this burden, should be sent to the Chief Information Officer, U.S. Patent and Trademark Office, u.S. Operatment of Commence, P. O. Box 1430, Alexandriu, V.S. 2213.1450. DO NOT SEND FEES OR COMPLETED FORMS TO THIS ADDRESS. SEND TO: Commissioner for Patents, P.O. Box 1450, Alexandria, V.A. 2213.1450.

## Privacy Act Statement

The Privacy Act of 1974 (P.L. 93-579) requires that you be given certain information in connection with your submission of the attached form related to a patient application or patient. Accordingly, pursuant to the requirements of the Act, please be advised that (1) the general authority for the collection of this information is \$3 U.S.C. 2(b)(2); (2) famishing of the information solicided is voluntary, and (3) the principal purpose for which the information is used by the U.S. Patient and Trademan Kollie is to process and/or examine your submission related to a patient application or patient. If you do not furnish the requested process and/or examine your submission related to a patient application or patient. If you do not furnish the requested related to the patient application or patient. If you do not furnish the requested related to the patient application or patient. If you do not furnish the requested related to the patient application or patient. If you do not furnish the requested to the patient process and/or examine your submission, which may related that the patient process and/or examine your submission, which may

The information provided by you in this form will be subject to the following routine uses:

- The information on this form will be treated confidentially to the extent allowed under the Freedom of Information Act (5 U.S.C. 552) and the Privacy Act (5 U.S.C. 552a). Records from this system of records may be disclosed to the Department of Justice to determine whether the Freedom of Information Act requires disclosure of these record s.
  - A record from this system of records may be disclosed, as a routine use, in the course of presenting evidence to a court, magistrate, or administrative tribunal, including disclosures to opposing counsel in the course of settlement neodiations.
  - A record in this system of records may be disclosed, as a routine use, to a Member of Congress submitting a request involving an individual, to whom the record partains, when the individual has requested assistance from the Member with respect to the subject matter of the record.
  - A record in this system of records may be disclosed, as a routine use, to a contractor of the Agency having need for the information in order to perform a contract. Recipients of information shall be required to comply with the requirements of the Privacy Act of 1974. as amended, pursuant to 5 U.S.C. 552a(m.).
  - A record related to an International Application filed under the Patent Cooperation Treaty in this system of records
    may be disclosed, as a routine use, to the International Bureau of the World Intellectual Property Organization, pursuant
    to the Patent Cooperation Treaty.
  - A record in this system of records may be disclosed, as a routine use, to another federal agency for purposes of National Security review (35 U.S.C. 181) and for review pursuant to the Atomic Energy Act (42 U.S.C. 218(c)).
  - 7. A record from this system of records may be disclosed, as a routine use, to the Administrator, General Services, or hisher designed, uturing an inspection of records conducted by GSA as part of that agency's responsibility to recommend improvements in records management practices and programs, under authority of 44 U.S.C. 2504 and 2506. Such disclosure shall be made in accordance with the GSA regulations governing inspection of records for this purpose, and any other relevant (i.e., GSA or Commerce) directive. Such disclosure shall not be used to make relevant (i.e., GSA or Commerce) directive. Such disclosure shall not be used to make relevant for the state of the s
- A record from this system of records may be disclosed, as a routine use, to the public after either publication of the application pursuant to 35 U.S. C. 12(b) or issuance of a patient pursuant to 35 U.S. C. 15.1 Further, a record may be disclosed, subject to the limitations of 37 CFR.114, as a routine use, to the public if the record via flori of mapplication which became abandoned or in which the proceedings were terminated and which application is referenced by either a published application, an application open to public inseptions or an issued patient.
  - A record from this system of records may be disclosed, as a routine use, to a Federal, State, or local law enforcement agency, if the USPTO becomes aware of a violation or potential violation of law or regulation.